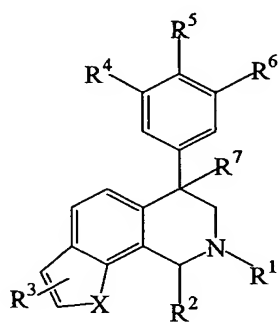


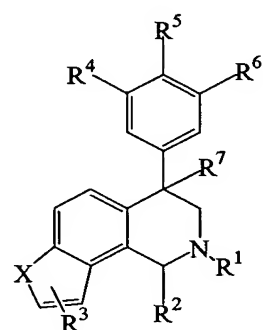
CLAIMS:

What is claimed is:

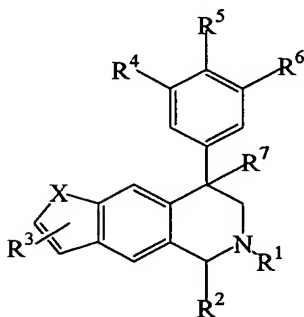
1. A method of treating chronic or neuropathic pain, treating or preventing
 5 migraine headache, or treating urge, stress or mixed urinary incontinence comprising
 administration of an effective amount of a compound selected from one of the
 Formulae IA, IB, IIA, IIB, IIIA or IIIB



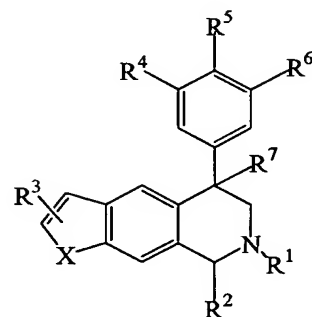
IA



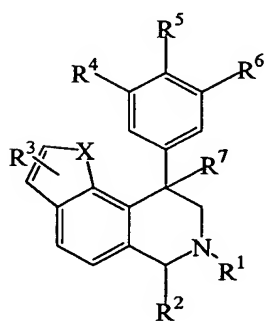
IB



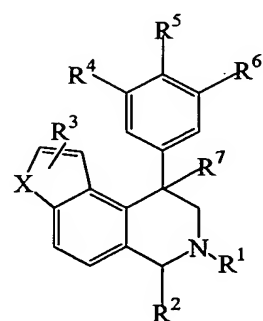
IIA



IIB



IIIA



IIIB

wherein:

R^1 is selected from the group consisting of C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, C_3 - C_6 cycloalkyl, C_4 - C_7 cycloalkylalkyl and benzyl, each of which is optionally substituted with 1 to 3 substituents independently selected at each occurrence from C_1 - C_3 alkyl, halogen, -CN, -OR⁸ and -NR⁸R⁹;

R^2 is selected from the group consisting of H, C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, C_3 - C_6 cycloalkyl, C_4 - C_7 cycloalkylalkyl and C_1 - C_6 haloalkyl;

R^3 is selected from the group consisting of H, halogen, C_1 - C_6 alkyl, C_1 - C_6 haloalkyl and C_3 - C_6 cycloalkyl, wherein C_1 - C_6 alkyl, C_1 - C_6 haloalkyl and C_3 - C_6 cycloalkyl are optionally substituted with 1 to 3 substituents selected independently at each occurrence from OR⁸ and NR⁸R⁹;

R^4 , R^5 , and R^6 are each independently selected at each occurrence thereof from the group consisting of H, halogen, -OR¹⁰, -NO₂, -NR¹⁰R¹¹, -NR¹⁰C(0)R¹¹, -NR¹⁰C(0)NR¹¹R¹², -S(0)_nR¹¹, -CN, -C(O)R¹¹, -C(O)₂R¹¹, -C(0)NR¹¹R¹², C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, C_3 - C_6 cycloalkyl and C_4 - C_7 cycloalkylalkyl, wherein each of C_1 - C_6 alkyl, C_2 - C_6 alkenyl, C_2 - C_6 alkynyl, C_3 - C_6 cycloalkyl and C_4 - C_7 cycloalkylalkyl are optionally substituted with 1 to 3 substituents independently selected at each occurrence with from C_1 - C_3 alkyl, halogen, =O, -CN, -OR⁸, -NR⁸R⁹ and phenyl, and wherein phenyl is optionally substituted 1-3 substituents selected independently at each occurrence from halogen, -CN, C_1 - C_4 alkyl, C_1 - C_4 haloalkyl, -OR⁸ and -NR⁸R⁹;

alternatively R^5 and R^6 taken together are -O-C(R¹¹)₂-O-;

R^7 is selected from the group consisting of H, halogen and OR¹⁰;

R^8 and R^9 are each independently selected from the group consisting of H, C_1 - C_4 alkyl, C_1 - C_4 haloalkyl, C_1 - C_4 alkoxyalkyl, C_1 - C_4 alkoxyalkylalkyl, C_3 - C_6 cycloalkyl, C_4 - C_7 cycloalkylalkyl, -C(0)R¹², phenyl and benzyl, wherein phenyl and benzyl are optionally substituted with 1 to 3 substituents selected independently at each

occurrence from halogen, cyano, C₁-C₄ alkyl, C₁-C₄ haloalkyl, C₁-C₄ alkoxy and C₁-C₄ haloalkoxy, or R⁸ and R⁹ are taken together with the nitrogen to which they are attached to form a piperidine, pyrrolidine, piperazine, N-methylpiperazine, morpholine, or thiomorpholine ring;

5

R¹⁰ is selected from the group consisting of H, C₁-C₄ alkyl, C₁-C₄ haloalkyl, C₁-C₄ alkoxyalkyl, C₃-C₆ cycloalkyl, C₄-C₇ cycloalkylalkyl, -C(O)R¹², phenyl and benzyl, wherein phenyl and benzyl are optionally substituted with 1 to 3 substituents selected independently at each occurrence from halogen, -NH₂, -OH, cyano, C₁-C₄ alkyl, C₁-C₄ haloalkyl, C₁-C₄ alkoxy and C₁-C₄ haloalkoxy;

10

R¹¹ is selected from the group consisting of H, C₁-C₄ alkyl, C₁-C₄ haloalkyl, C₁-C₄ alkoxyalkyl, C₃-C₆ cycloalkyl, C₄-C₇ cycloalkylalkyl, phenyl and benzyl, where phenyl and benzyl are optionally substituted with 1 to 3 substituents selected independently at each occurrence from halogen, -NH₂, -OH, cyano, C₁-C₄ alkyl, C₁-C₄ haloalkyl, C₁-C₄ alkoxy and C₁-C₄ haloalkoxy, or R¹⁰ and R¹¹ are taken together with the nitrogen to which they are attached to form a piperidine, pyrrolidine, N-methylpiperazine, morpholine, or thiomorpholine ring, with the proviso that only one of R⁸ and R⁹ or R¹⁰ and R¹¹ are taken together with the nitrogen to which they are attached to form a piperidine, pyrrolidine, piperazine, N-methylpiperazine, morpholine, or thiomorpholine ring;

15

20

R¹² is selected from the group consisting of C₁-C₄ alkyl, C₁-C₄ haloalkyl and phenyl;

25 X is selected from the group consisting of 0, NR¹³ and S;

the ring containing X is selected from furan, pyrrole, thiophene, dihydrofuran, dihydropyrrole, and dihydrothiophene; n is 0, 1, or 2; and,

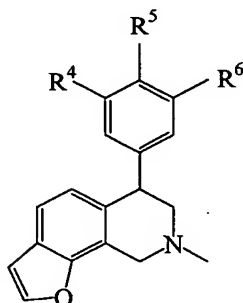
30 R¹³ is selected from the group consisting of H, C₁-C₆ alkyl, benzyl and phenyl, wherein C₁-C₆ alkyl, benzyl and phenyl are optionally substituted with 1-3 substituents independently at each occurrence from halogen, -NH₂, -OH, cyano, C₁-C₄ alkyl, C₁-C₄ haloalkyl, C₁-C₄ alkoxy and C₁-C₄ haloalkoxy;

or a pharmaceutically acceptable salt thereof or an isomer or prodrug thereof to a patient in need thereof.

- 5 2. A method of claim 1, wherein R¹ is C₁-C₆ alkyl.
3. A method of claim 2, wherein R¹ is CH₃.
4. A method of claim 1, wherein R² is H, C₁-C₆ alkyl, C₃-C₆ cycloalkyl, or C₁-C₆
10 haloalkyl.
5. A method of claim 4, wherein R² is H or C₁-C₆ alkyl.
6. A method of claim 5, wherein R² is H.
- 15 7. A method of claim 1, wherein R³ is at each occurrence thereof independently
H, halogen, C₁-C₆ alkyl, or C₁-C₆ alkyl substituted with from 1 to 3 of OR⁸ or NR⁸R⁹.
8. A method of claim 7, wherein R³ is H or C₁-C₆ alkyl.
- 20 9. A method of claim 8, wherein R³ is H.
10. A method of claim 1, wherein R¹ is CH₃, R² is H and R³ is H.
- 25 11. A method of claim 1, wherein R⁴, R⁵ and R⁶ are each independently H, halogen,
C₁-C₆ alkyl or -OR¹⁰.
12. A method of claim 11, wherein at least one of R⁴, R⁵ and R⁶ is H.
- 30 13. A method of claim 12, wherein each of R⁴, R⁵ and R⁶ are H.
14. A method of claim 12, wherein one of R⁴, R⁵ and R⁶ is halogen.

15. A method of claim 1, wherein R^1 is CH_3 , R^2 and R^3 are each H, and at least one of R^4 , R^5 , and R^6 is H.

16. A method of claim 1 wherein the compound is a compound of Formula (10):



(10)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

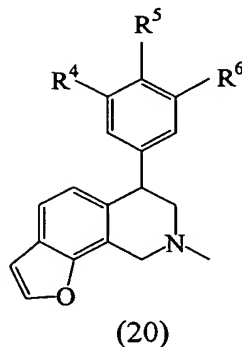
a compound of Formula (10) wherein R^4 is H, R^5 is H and R^6 is H;

a compound of Formula (10) wherein R^4 is H, R^5 is Me and R^6 is H;

a compound of Formula (10) wherein R^4 is Cl, R^5 is H and R^6 is H; and

a compound of Formula (10) wherein R^4 is H, R^5 is F and R^6 is H.

17. A method of claim 1 wherein the compound is a compound of Formula (20):



or a pharmaceutically acceptable salt form thereof selected from the group consisting
5 essentially of:

a compound of Formula (20) wherein R^4 is H, R^5 is H and R^6 is H;

a compound of Formula (20) wherein R^4 is H, R^5 is Me and R^6 is H;

10

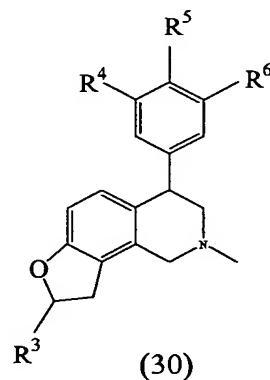
a compound of Formula (20) wherein R^4 is H, R^5 is Cl and R^6 is H;

a compound of Formula (20) wherein R^4 is H, R^5 is F and R^6 is H; and

15

a compound of Formula (20) wherein R^4 is F, R^5 is H and R^6 is F.

18. A method of claim 1 wherein the compound is a compound of Formula (30):



or a pharmaceutically acceptable salt form thereof selected from the group consisting
20 essentially of:

a compound of Formula (30) wherein R^3 is H, R^4 is H, R^5 is H and R^6 is H;

a compound of Formula (30) wherein R^3 is H, R^4 is F, R^5 is F and R^6 is H;

5 a compound of Formula (30) wherein R^3 is H, R^4 is F, R^5 is H and R^6 is F;

a compound of Formula (30) wherein R^3 is H, R^4 is H, R^5 is F and R^6 is H;

10 a compound of Formula (30) wherein R^3 is H, R^4 is Cl, R^5 is H and R^6 is H;

a compound of Formula (30) wherein R^3 is H, R^4 is H, R^5 is Cl and R^6 is H;

a compound of Formula (30) wherein R^3 is H, R^4 is H, R^5 is Cl and R^6 is F;

15 a compound of Formula (30) wherein R^3 is H, R^4 is H, R^5 is F and R^6 is Cl;

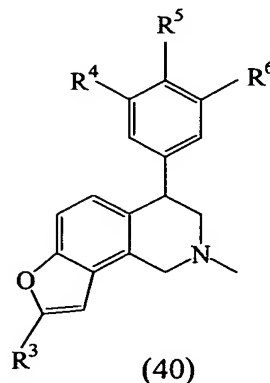
a compound of Formula (30) wherein R^3 is H, R^4 is F, R^5 is H and R^6 is Cl;

a compound of Formula (30) wherein R^3 is H, R^4 is H, R^5 is OMe and R^6 is H;

20 and

a compound of Formula (30) wherein R^3 is H, R^4 is F, R^5 is H and R^6 is H.

19. A method of claim 1 wherein the compound is a compound of Formula (40):

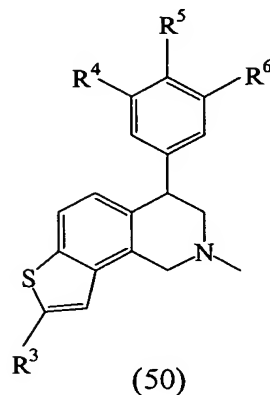


25

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

- 5 a compound of Formula (40) wherein R^3 is H, R^4 is H, R^5 is H and R^6 is H;
- a compound of Formula (40) wherein R^3 is H, R^4 is F, R^5 is F and R^6 is H;
- a compound of Formula (40) wherein R^3 is H, R^4 is F, R^5 is H and R^6 is F;
- 10 a compound of Formula (40) wherein R^3 is H, R^4 is F, R^5 is H and R^6 is H;
- a compound of Formula (40) wherein R^3 is H, R^4 is H, R^5 is F and R^6 is H;
- a compound of Formula (40) wherein R^3 is H, R^4 is Cl, R^5 is H and R^6 is H;
- 15 a compound of Formula (40) wherein R^3 is H, R^4 is H, R^5 is Cl and R^6 is H;
- a compound of Formula (40) wherein R^3 is H, R^4 is H, R^5 is Cl and R^6 is F;
- 20 a compound of Formula (40) wherein R^3 is H, R^4 is H, R^5 is F and R^6 is Cl;
- a compound of Formula (40) wherein R^3 is H, R^4 is F, R^5 is H and R^6 is Cl;
- a compound of Formula (40) wherein R^3 is H, R^4 is H, R^5 is OMe and R^6 is H;
- 25 a compound of Formula (40) wherein R^3 is Me, R^4 is H, R^5 is H and R^6 is H;
- a compound of Formula (40) wherein R^3 is Et, R^4 is H, R^5 is H and R^6 is H;
- and
- 30 a compound of Formula (40) wherein R^3 is CH_2OH , R^4 is H, R^5 is H and R^6 is H.

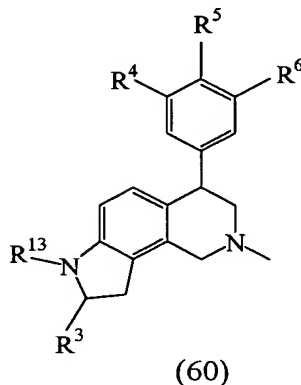
20. A method of claim 1 wherein the compound is a compound of Formula (50):



or a pharmaceutically acceptable salt form thereof selected from the group consisting
5 essentially of:

a compound of Formula (50) wherein R^3 is H, R^4 is H, R^5 is H and R^6 is H.

21. A method of claim 1 wherein the compound is a compound of Formula (60):



10

or a pharmaceutically acceptable salt form thereof selected from the group consisting
essentially of:

15 a compound of Formula (60) wherein R^3 is H, R^4 is H, R^5 is H, R^6 is H and R^{13}
is H;

a compound of Formula (60) wherein R^3 is H, R^4 is H, R^5 is H, R^6 is H and R^{13}
is Me;

20

a compound of Formula (60) wherein R^3 is H, R^4 is H, R^5 is H, R^6 is H and R^{13} is Et;

5 a compound of Formula (60) wherein R^3 is H, R^4 is H, R^5 is F, R^6 is F and R^{13} is H;

a compound of Formula (60) wherein R^3 is H, R^4 is H, R^5 is F, R^6 is F and R^{13} is Me;

10 a compound of Formula (60) wherein R^3 is H, R^4 is F, R^5 is H, R^6 is F and R^{13} is H;

a compound of Formula (60) wherein R^3 is H, R^4 is F, R^5 is H, R^6 is F and R^{13} is Me;

15 a compound of Formula (60) wherein R^3 is H, R^4 is Cl, R^5 is H, R^6 is H and R^{13} is H;

a compound of Formula (60) wherein R^3 is H, R^4 is Cl, R^5 is H, R^6 is H and R^{13} is Me;

20 a compound of Formula (60) wherein R^3 is H, R^4 is F, R^5 is H, R^6 is H and R^{13} is H;

25 a compound of Formula (60) wherein R^3 is H, R^4 is H, R^5 is F, R^6 is H and R^{13} is H;

a compound of Formula (60) wherein R^3 is H, R^4 is F, R^5 is Cl, R^6 is H and R^{13} is H;

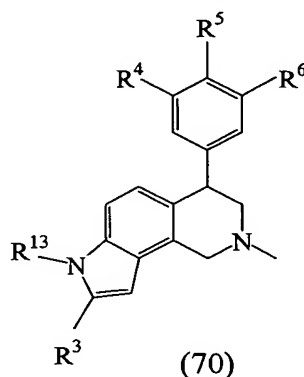
30 a compound of Formula (60) wherein R^3 is H, R^4 is F, R^5 is Cl, R^6 is H and R^{13} is Me;

a compound of Formula (60) wherein R^3 is H, R^4 is Cl, R^5 is F, R^6 is H and R^{13} is H; and

a compound of Formula (60) wherein R^3 is H, R^4 is Cl, R^5 is F, R^6 is H and R^{13} is Me.

5

22. A method of claim 1 wherein the compound is a compound of Formula (70):



or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

10

a compound of Formula (70) wherein R^3 is H, R^4 is H, R^5 is H, R^6 is H and R^{13} is H;

15 a compound of Formula (70) wherein R^3 is H, R^4 is H, R^5 is H, R^6 is H and R^{13} is Me;

a compound of Formula (70) wherein R^3 is H, R^4 is H, R^5 is H, R^6 is H and R^{13} is Et;

20

a compound of Formula (70) wherein R^3 is H, R^4 is H, R^5 is H, R^6 is H and R^{13} is Bn;

25 a compound of Formula (70) wherein R^3 is H, R^4 is H, R^5 is F, R^6 is F and R^{13} is H;

a compound of Formula (70) wherein R^3 is H, R^4 is H, R^5 is F, R^6 is F and R^{13} is Me;

5 a compound of Formula (70) wherein R^3 is H, R^4 is F, R^5 is H, R^6 is F and R^{13} is Me;

a compound of Formula (70) wherein R^3 is H, R^4 is Cl, R^5 is H, R^6 is H and R^{13} is H;

10 a compound of Formula (70) wherein R^3 is H, R^4 is Cl, R^5 is H, R^6 is H and R^{13} is Me;

a compound of Formula (70) wherein R^3 is H, R^4 is F, R^5 is H, R^6 is H and R^{13} is H;

15 a compound of Formula (70) wherein R^3 is H, R^4 is F, R^5 is H, R^6 is H and R^{13} is Me;

a compound of Formula (70) wherein R^3 is H, R^4 is H, R^5 is F, R^6 is H and R^{13} is H;

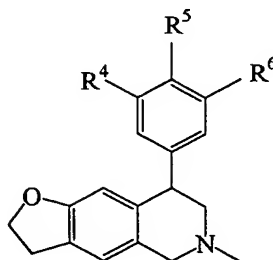
20 a compound of Formula (70) wherein R^3 is H, R^4 is F, R^5 is Cl, R^6 is H and R^{13} is H;

25 a compound of Formula (70) wherein R^3 is H, R^4 is F, R^5 is Cl, R^6 is H and R^{13} is Me;

a compound of Formula (70) wherein R^3 is H, R^4 is Cl, R^5 is F, R^6 is H and R^{13} is H; and

30 a compound of Formula (70) wherein R^3 is H, R^4 is Cl, R^5 is F, R^6 is H and R^{13} is Me.

23. A method of claim 1 wherein the compound is a compound of Formula (80):



(80)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

5

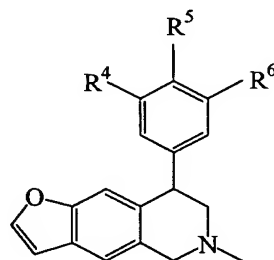
a compound of Formula (80) wherein R⁴ is H, R⁵ is H and R⁶ is H;

a compound of Formula (80) wherein R⁴ is H, R⁵ is F and R⁶ is H; and

10

a compound of Formula (80) wherein R⁴ is H, R⁵ is F and R⁶ is F.

24. A method of claim 1 wherein the compound is a compound of Formula (90):



(90)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

15

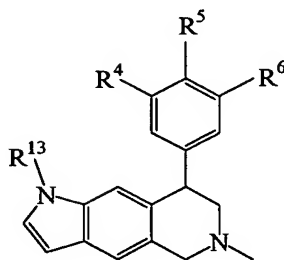
a compound of Formula (90) wherein R⁴ is H, R⁵ is H and R⁶ is H;

a compound of Formula (90) wherein R⁴ is H, R⁵ is F and R⁶ is F; and

20

a compound of Formula (90) wherein R⁴ is H, R⁵ is F and R⁶ is H.

25. A method of claim 1 wherein the compound is a compound of Formula (100):



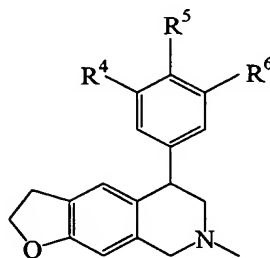
(100)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

5

a compound of Formula (100) wherein R^4 is H, R^5 is H, R^6 is H and R^{13} is H.

26. A method of claim 1 wherein the compound is a compound of Formula (110):



(110)

10 or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (110) wherein R^4 is H, R^5 is H and R^6 is H;

15

a compound of Formula (110) wherein R^4 is H, R^5 is F and R^6 is F;

a compound of Formula (110) wherein R^4 is H, R^5 is F and R^6 is H;

a compound of Formula (110) wherein R^4 is H, R^5 is H and R^6 is Cl;

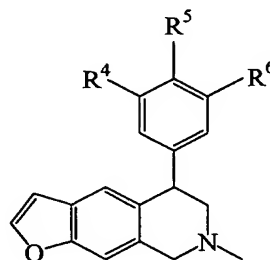
20

a compound of Formula (110) wherein R^4 is H, R^5 is Cl and R^6 is F;

a compound of Formula (110) wherein R^4 is H, R^5 is F and R^6 is Cl; and

a compound of Formula (110) wherein R^4 is H, R^5 is OMe and R^6 is H.

- 5 27. A method of claim 1 wherein the compound is a compound of Formula (120):



(120)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

- 10 a compound of Formula (120) wherein R^4 is H, R^5 is H and R^6 is H;

a compound of Formula (120) wherein R^4 is H, R^5 is F and R^6 is F;

a compound of Formula (120) wherein R^4 is H, R^5 is F and R^6 is H;

15

a compound of Formula (120) wherein R^4 is H, R^5 is H and R^6 is Cl;

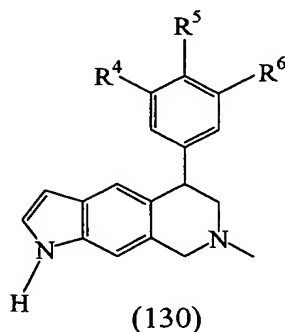
a compound of Formula (120) wherein R^4 is H, R^5 is Cl and R^6 is F;

- 20 a compound of Formula (120) wherein R^4 is H, R^5 is OMe and R^6 is H; and

a compound of Formula (120) wherein R^4 is H, R^5 is F and R^6 is Cl.

25

28. A method of claim 1 wherein the compound is a compound of Formula (130):

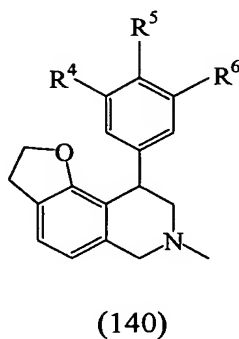


or a pharmaceutically acceptable salt form thereof selected from the group consisting
essentially of:

a compound of Formula (130) wherein R⁴ is H, R⁵ is H and R⁶ is H; and

a compound of Formula (130) wherein R⁴ is H, R⁵ is Bn and R⁶ is H.

29. A method of claim 1 wherein the compound is a compound of Formula (140):



or a pharmaceutically acceptable salt form thereof selected from the group consisting
essentially of:

a compound of Formula (140) wherein R⁴ is H, R⁵ is H and R⁶ is H;

a compound of Formula (140) wherein R⁴ is H, R⁵ is F and R⁶ is H;

a compound of Formula (140) wherein R⁴ is H, R⁵ is F and R⁶ is Cl;

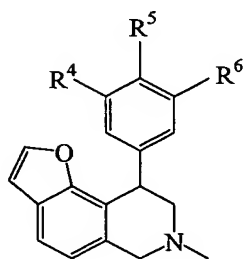
a compound of Formula (140) wherein R^4 is H, R^5 is Cl and R^6 is F;

a compound of Formula (140) wherein R^4 is H, R^5 is H and R^6 is Cl;

5 a compound of Formula (140) wherein R^4 is H, R^5 is OMe and R^6 is H;

a compound of Formula (140) wherein R^4 is H, R^5 is F and R^6 is F.

30. A method of claim 1 wherein the compound is a compound of Formula (150):



(150)

10 or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (150) wherein R^4 is H, R^5 is H and R^6 is H;

15 a compound of Formula (150) wherein R^4 is H, R^5 is F and R^6 is H;

a compound of Formula (150) wherein R^4 is H, R^5 is F and R^6 is Cl;

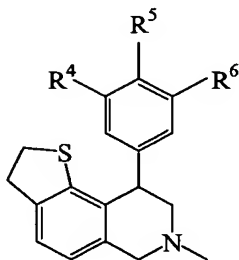
20 a compound of Formula (150) wherein R^4 is H, R^5 is Cl and R^6 is F;

a compound of Formula (150) wherein R^4 is H, R^5 is H and R^6 is Cl;

a compound of Formula (150) wherein R^4 is H, R^5 is OMe and R^6 is H; and

25 a compound of Formula (150) wherein R^4 is H, R^5 is F and R^6 is F.

31. A method of claim 1 wherein the compound is a compound of Formula (160):



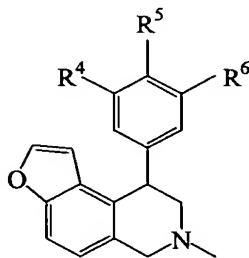
(160)

or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

5

a compound of Formula (160) wherein R^4 is H, R^5 is H and R^6 is H.

32. A method of claim 1 wherein the compound is a compound of Formula (170):



(170)

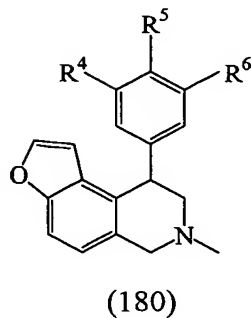
10 or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

a compound of Formula (170) wherein R^4 is H, R^5 is H and R^6 is H;

15 a compound of Formula (170) wherein R^4 is H, R^5 is F and R^6 is H; and

a compound of Formula (170) wherein R^4 is H, R^5 is F and R^6 is F.

33. A method of claim 1 wherein the compound is a compound of Formula (180):



or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

5

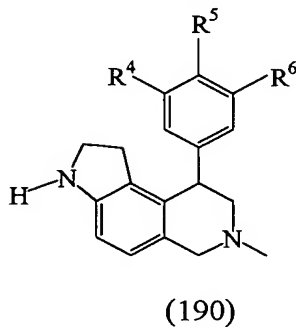
a compound of Formula (180) wherein R^4 is H, R^5 is H and R^6 is H;

a compound of Formula (180) wherein R^4 is H, R^5 is F and R^6 is H; and

10

a compound of Formula (180) wherein R^4 is H, R^5 is F and R^6 is F.

34. A method of claim 1 wherein the compound is a compound of Formula (190):

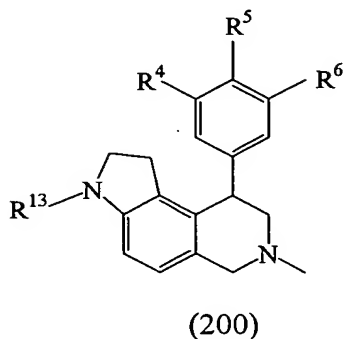


or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

15

a compound of Formula (190) wherein R^4 is H, R^5 is H and R^6 is H.

35. A method of claim 1 wherein the compound is a compound of Formula (200):



or a pharmaceutically acceptable salt form thereof selected from the group consisting essentially of:

5

a compound of Formula (200) wherein R⁴ is H, R⁵ is H, R⁶ is H and R¹³ is H;
and

10

a compound of Formula (200) wherein R⁴ is H, R⁵ is H, R⁶ is H and R¹³ is Me.

36. A method of claim 1 wherein the compound is selected from the group consisting of:

15

(R)-2-methyl-4-phenyl-1,2,3,4,8,9-hexahydro-furo[2,3-h]isoquinoline;

(S)-2-methyl-4-phenyl-1,2,3,4,8,9-hexahydro-furo[2, 3-h]isoquinoline;

(R)-7-methyl-5-phenyl-5,6,7,8-tetrahydro-furo[3,2-g]isoquinoline;

20

(S)-7-methyl-5-phenyl-5,6,7,8-tetrahydro-furo[3, 2-g]isoquinoline;

(R)-4-(4-fluoro-phenyl)-2-methyl-1,2,3,4- tetrahydrofuro[2,3-h]isoquinoline;

(S)-4-(4-fluoro-phenyl)-2-methyl-1,2,3,4-tetrahydrofuro[2,3-h]isoquinoline;

25

(R)-4-(3,4-difluoro-phenyl)-2-methyl-1,2,3,4-tetrahydrofuro[2, 3-h]isoquinoline;

(S)-4-(3, 4-difluoro-phenyl)-2-methyl-1,2,3,4-tetrahydrofuro[2,3-h]isoquinoline;

(R)-2-methyl-4-phenyl-1,2,3,4-tetrahydro-furo[2, 3-h]isoquinoline;

5

(S)-2-methyl-4-phenyl-1,2,3,4-tetrahydro-furo[2, 3-h]isoquinoline;

(R)-4-(4-chloro-phenyl)-2-methyl-1,2,3,4- tetrahydro-furo[2, 3-h]isoquinoline;

10

(S)-4-(4-chloro-phenyl)-2- methyl-1,2,3,4-tetrahydrofuro[2,3- h]isoquinoline;

(R)-8-methyl- 6-phenyl-2,3,6,7,8,9-hexahydro-furo[3,2-h]isoquinoline;

(S)-8-methyl-6-phenyl-2,3,6,7,8,9-hexahydro-furo[3,2-h]isoquinoline;

15

(R)-4-(4-fluoro-phenyl)-2-methyl-1,2,3,4-tetrahydrofuro[2,3-h]isoquinoline;

(S)-4-(4-fluoro-phenyl)-2-methyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;

20

(R)-4-(3,5-difluoro-phenyl)-2-methyl-1,2,3, 4-tetrahydrofuro[2,3-h]isoquinoline;

(S)-4-(3,5-difluoro-phenyl)-2-methyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;

25

(R)-2-methyl-4-phenyl2,3,4,7-tetrahydro-1H-pyrrolo[2,3-h]isoquinoline; and

(S)-2-methyl-4-phenyl-2,3,4,7-tetrahydro-1H-pyrrolo[2, 3-h]isoquinoline.

30

37. A method of claim 1 wherein the compound is selected from the group consisting of:

(+)-2-methyl-4-phenyl-1,2,3,4,8,9-hexahydro-furo[2, 3-h]isoquinoline;

- (-)-2-methyl-4-phenyl-1,2,3,4,8,9-hexahydro-furo[2,3-h]isoquinoline;
- (+)-7-methyl-5-phenyl-5,6,7,8-tetrahydro-furo[3, 2-g]isoquinoline;
- 5 (-)-7-methyl-5-phenyl-5,6,7,8-tetrahydro-furo[3, 2-g]isoquinoline;
- (+)-4-(4-fluoro-phenyl)-2-methyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;
- 10 (-)-4-(4-fluoro-phenyl)-2-methyl-1,2,3,4- tetrahydro-furo[2,3-h]isoquinoline;
- (+)-4-(3,4-difluoro-phenyl)-2-methyl-1,2,3,4-tetrahydro-furo[2,3-
h]isoquinoline;
- 15 (-)-4-(3,4- difluoro-phenyl)-2-methyl-1,2,3,4-tetrahydro-furo[2,3-
h]isoquinoline;
- (+)-2-methyl-4-phenyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;
- 20 (-)-2-methyl-4-phenyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;
- (+)-4-(4-chloro-phenyl)-2-methyl-1,2,3,4- tetrahydrofuro[2,3-h]isoquinoline;
- (-)-4-(4-chloro-phenyl)-2-methyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;
- 25 (+)-8-methyl-6-phenyl-2,3,6,7,8,9-hexahydro-furo[3,2h]isoquinoline;
- (-)-8-methyl- 6-phenyl-2,3,6,7,8,9-hexahydro-furo[3,2-h]isoquinoline;
- 30 (+)-4-(4- fluoro-phenyl)-2-methyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;
- (-)-4-(4-fluoro-phenyl)-2-methyl-1,2,3,4- tetrahydrofuro[2,3-h]isoquinoline;

(+)-4-(3,5-difluoro-phenyl)-2-methyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;

5 (-)-4-(3,5-difluoro-phenyl)-2-methyl-1,2,3,4-tetrahydro-furo[2,3-h]isoquinoline;

(+)-2-methyl-4-phenyl-2,3,4,7-tetrahydro-1H-pyrrolo[2,3-h]isoquinoline; and

(-)-2-methyl-4-phenyl-2,3,4,7-tetrahydro-1H- pyrrolo[2,3-h]isoquinoline.

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